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# **Risk Adjustment Regulations Draft for Public Comment (VOTE)**

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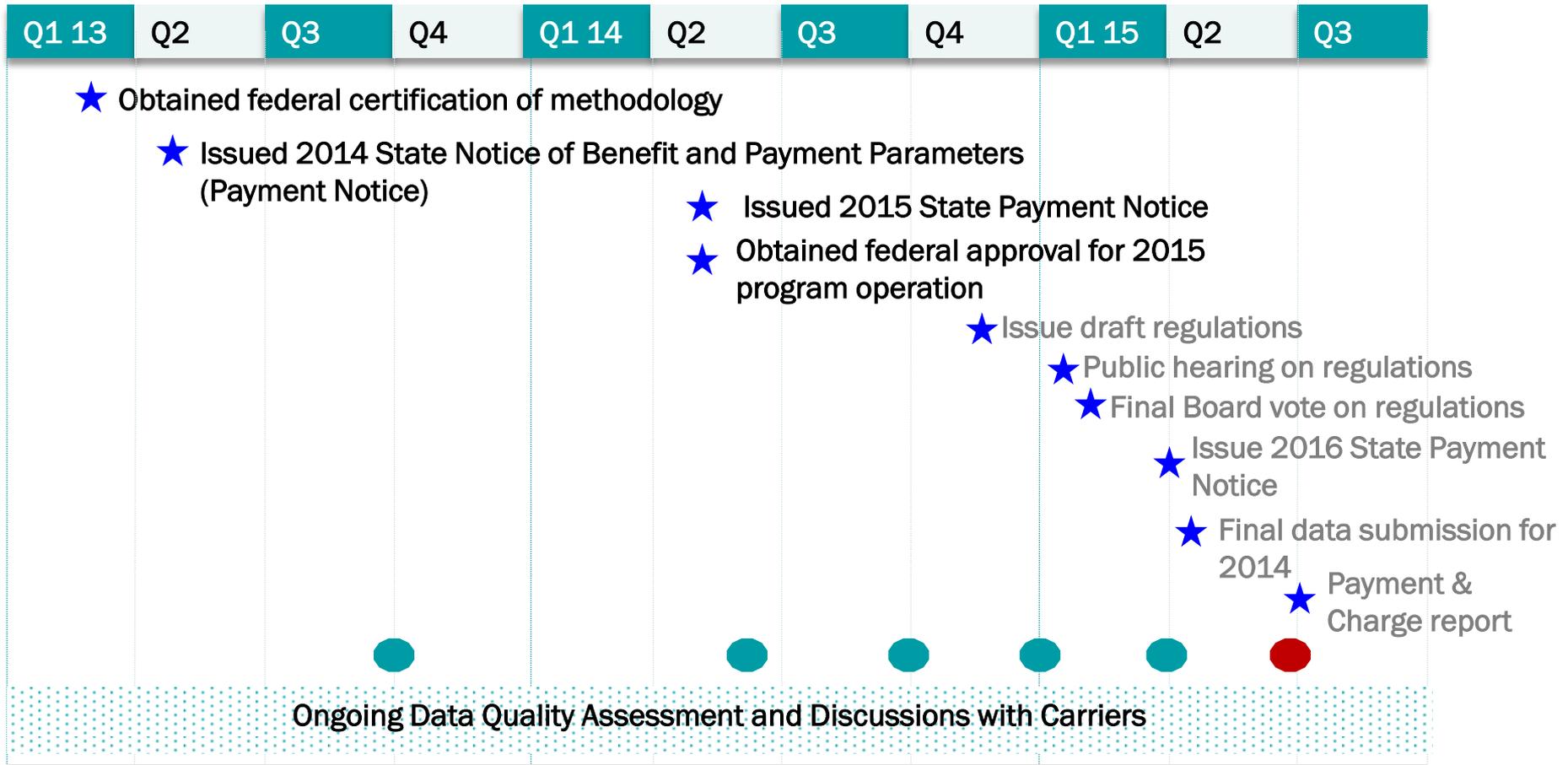
Board of Directors Meeting, December 11, 2014

# Background



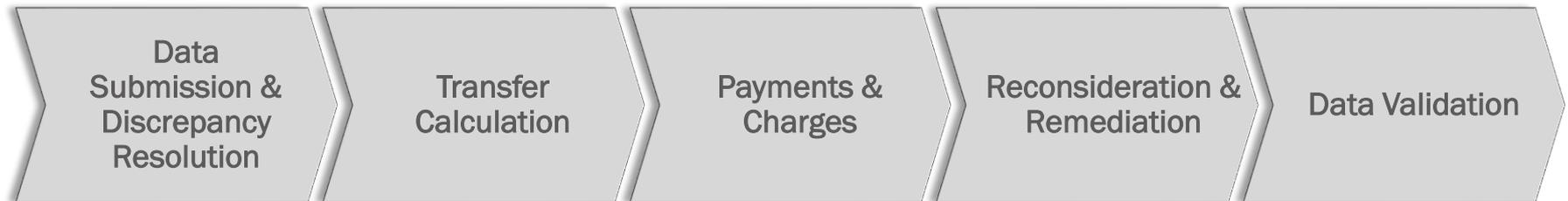
- The ACA requires the implementation of a risk adjustment program for the individual and small group market, effective 2014. The objective of the program is to help maintain a level-playing field, stabilize premium and promote competition
- States have the option to administer state-based programs or default to the federal program. In consultation with issuers and other stakeholders, Massachusetts pursued a state-based approach and successfully obtained the necessary federal to operate the program
- Significant progress has been made in risk adjustment implementation over the past two years, including establishing key steps in data collection, data quality improvement, program simulation and reporting
- Today we are proposing to issue draft Risk Adjustment regulations for public comment. By ultimately promulgating these regulations, we intend to formalize a series of program rules that are essential to the reliable operationalization of Risk Adjustment, in advance of the upcoming 2014 Benefit Year payment and charge settlement

# Key Milestones



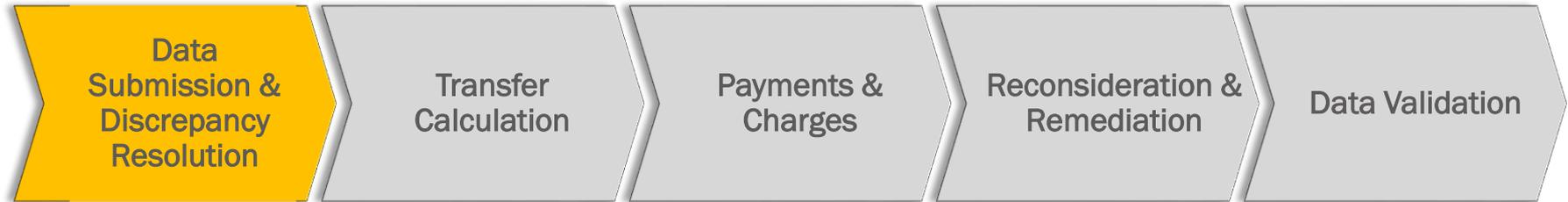
- Market-wide Simulation
- Funds Settlement Calculations for 2014

# Summary of Risk Adjustment Processes



- Risk adjustment is a multi-phase process
- Our proposed regulations closely align with United States Department of Health and Human Services (HHS) approach

# Data Submission & Discrepancy Resolution



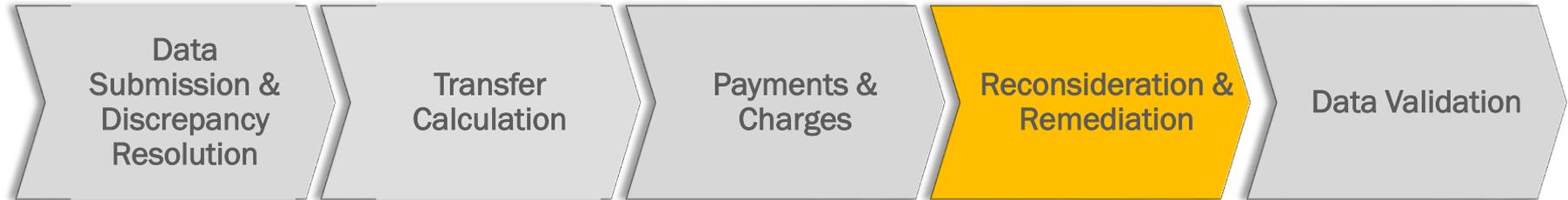
- Data submission requirements
  - Issuers must comply with the Center for Health Information and Analysis (CHIA) data submission requirements, including data privacy & security requirements
  - All data for a given benefit year must be submitted by April 30th of following calendar year
- Carriers review reports, identify discrepancies
  - Member month tracker reports will be provided to issuers each month, and simulation reports each quarter
  - Within 30 days, issuers must either: 1) confirm accuracy of report or 2) identify discrepancies
  - The Health Connector will work with issuers to resolve discrepancies within 60 days or close cases at its discretion
- April 2015 – Final discrepancy identification & resolution period
  - By March 2015, the Health Connector will provide a quarterly simulation report representing data from the 1st three quarters of 2014, as well as the final member month tracker report for the entire benefit year
  - All data must be locked down by April 30th; any discrepancies raised but not resolved by this date, or discrepancies that could not have been raised will be addressed through the reconsideration process

# Transfer Calculations, Payments & Charges



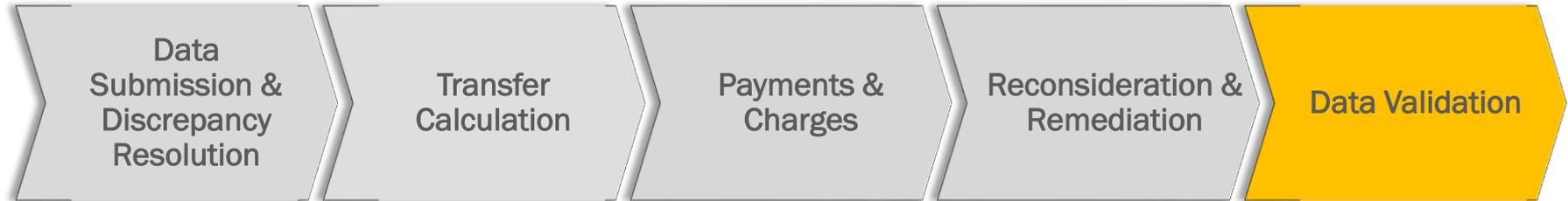
- Transfer calculations
  - May-June: Health Connector to calculate risk scores and determine payments and charges
  - By July 1st: Health Connector to send payment/charge report to issuers
  - Calculation will be based on data finalized as of April 30
  - Issuers who do not submit data on time may be subject to default charge
- Issuers who owe charges
  - Issuers who owe charges will have 30 days from the date the report is issued to submit charges
  - Late payments will be assessed interest at 12% per annum, which begins accruing 40 days from the date of the final payment/charge report
  - Payment and interest may not be stayed pending reconsideration – any adjustments that arise from the reconsideration process will be applied on a prospective basis
- Payments to recipient issuers
  - Transfer payments will be made after and only to the extent of receipt of risk adjustment charges
  - If the Health Connector does not receive full submission of charges owed on time, payments to recipient issuers will be made on a pro-rated basis, with additional distribution cycles in line with when the Health Connector receives subsequent submissions

# Reconsideration & Remedies



- Grounds for a request for reconsideration
  - Issuers may request reconsideration with respect to: 1) incorrect application of the methodology (including unresolved data discrepancies), or 2) mathematical errors. The methodology itself is not subject to request for reconsideration
  - Materiality threshold: dispute amount no less than 1% of the applicable payment/charges or \$25,000, whichever is less
- Reconsideration request
  - Issuers must file written requests for reconsideration within 30 days of the payment/charge report, and must specify the reasons for the challenge and the amount in dispute
  - Issuers may submit documentary evidence – but may not submit evidence that could have been submitted during discrepancy resolution
- Review
  - First-level review – reconsideration by the Health Connector
  - Second-level review – reconsideration through hearing, conducted according to the Massachusetts procedures/policies for informal hearings and any administrative bulletins issued by the Health Connector
- Remediation
  - Any change to a payment/charge resulting from the reconsideration process will be applied prospectively to future years' risk adjustment transfers after issuance of the decision and conclusion of any proceedings for judicial review

# Data Validation



- Risk Adjustment Data Validation (RADV) is a retrospective process of examining the quality of risk adjustment data, for purpose of validating and potentially modifying risk adjustment settlement as appropriate
- In the 2014 State Notice of Benefit and Payment Parameters, we described three different options for evaluation: 1) statistic-driven approach with limited medical record review; 2) single-level medical review; and 3) two-level medical record review
- Key considerations include: credibility of methodology; availability of vendors and tools; administrative feasibility; etc.
- Have been in regular discussion with carriers as well as the federal Centers for Medicare and Medicaid Services (CMS)
  - RADV is not part of the risk adjustment methodology subject to federal certification; state has more flexibility in defining and refining processes overtime, although RADV is still considered in CMS's operational approval
- Consistent with HHS approach, RADV for 2014 will be for informational purposes only, without actual adjustment of payments and charges

# Regulation Timeline



- Issuance of draft regulations will begin period of public comment
- Public hearing is scheduled for January 28, 2015
- Following public hearing and close of public comment, draft regulations will be amended and finalized for a final vote by this Board at the February 12, 2015 meeting

# Vote Language

**Moved that the Board issue regulations regarding risk adjustment, 956 CMR 13.00, in draft form for public hearing and comment.**